

**Amendments to the Claims**

Please amend Claims 1, 2, 5, 6 and 9. Please add new Claims 13-28. The Claim Listing below will replace all prior versions of the claims in the application:

**Claim Listing**

1. (Currently Amended) A method for treating a condition characterized by activation of the inflammatory cytokine cascade, comprising administering an ~~effective~~ amount of an antagonist or inhibitor of HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.
2. (Currently Amended) The method of Claim 1 further comprising administering a second agent in combination with the ~~antagonist or inhibitor of HMG1~~ antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
3. (Original) The method of Claim 2 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
4. (Original) The method of Claim 3 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
5. (Currently Amended) A method for treating sepsis ~~and related conditions involving activation of the inflammatory cytokine cascade~~, comprising administering an ~~effective~~ amount of an ~~antagonist or inhibitor of HMG1~~ antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.

6. (Currently amended) The method of Claim 5 further comprising administering a second agent in combination with the ~~antagonist or inhibitor of HMG1~~ antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
7. (Original) The method of Claim 6 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
8. (Original) The method of Claim 7 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
9. (Currently Amended) A method for treating rheumatoid arthritis, comprising administering an ~~effective amount of an~~ antagonist or inhibitor of HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.
10. (Original) The method of Claim 9 further comprising administering a second agent in combination with the HMG1 antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
11. (Original) The method of Claim 10 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
12. (Original) The method of Claim 11 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
13. (New) A method for treating inflammatory bowel disease, comprising administering an amount of an HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.

14. (New) The method of Claim 13 further comprising administering a second agent in combination with the HMG1 antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
15. (New) The method of Claim 14 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
16. (New) The method of Claim 15 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
17. (New) A method for treating systemic lupus erythematosus, comprising administering an amount of an HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.
18. (New) The method of Claim 17 further comprising administering a second agent in combination with the HMG1 antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
19. (New) The method of Claim 18 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
20. (New) The method of Claim 19 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
21. (New) A method for treating psoriasis, comprising administering an amount of an HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.

22. (New) The method of Claim 21 further comprising administering a second agent in combination with the HMG1 antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
23. (New) The method of Claim 22 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
24. (New) The method of Claim 23 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).
25. (New) A method for treating cardiovascular disease, comprising administering an amount of an HMG1 antagonist effective to inhibit the inflammatory cytokine cascade, wherein said HMG1 antagonist is an antibody that binds to HMG1 and inhibits the interaction between HMG1 and RAGE.
26. (New) The method of Claim 25 further comprising administering a second agent in combination with the HMG1 antagonist, wherein the second agent is an antagonist of an early sepsis mediator.
27. (New) The method of Claim 26 wherein the second agent is an antagonist of a cytokine selected from the group consisting of TNF, IL-1 $\alpha$ , IL-1 $\beta$ , MIF and IL-6.
28. (New) The method of Claim 27 wherein the second agent is an antibody to TNF or an IL-1 receptor antagonist (IL-1ra).